PIGA gene

phosphatidylinositol glycan anchor biosynthesis class A

Normal Function

The *PIGA* gene provides instructions for making a protein called phosphatidylinositol glycan class A. This protein takes part in a series of steps that produce a molecule called GPI anchor. Specifically, phosphatidylinositol glycan class A is involved in the first step of the sequence, which produces an intermediate molecule called N-acetylglucosaminyl phosphatidylinositol, or GlcNAc-PI. This step takes place in the endoplasmic reticulum of the cell, a structure involved in protein processing and transport. The PIGA protein forms a complex with several other proteins, and this complex helps to start the reaction that produces GlcNAc-PI.

The GPI anchor, the ultimate product of the sequence, attaches many different proteins to the cell membrane, thereby ensuring that these proteins are available when needed at the surface of the cell.

Health Conditions Related to Genetic Changes

paroxysmal nocturnal hemoglobinuria

Some gene mutations are acquired during a person's lifetime and are present only in certain cells. These changes, which are called somatic mutations, are not inherited. In people with paroxysmal nocturnal hemoglobinuria, somatic mutations of the *PIGA* gene occur in blood-forming cells called hematopoietic stem cells. Hematopoietic stem cells produce red blood cells (erythrocytes) that carry oxygen, white blood cells (leukocytes) that protect the body from infection, and platelets (thrombocytes) that are involved in blood clotting.

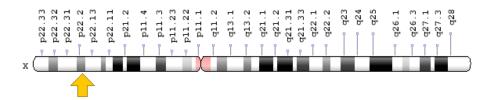
Individuals with paroxysmal nocturnal hemoglobinuria have one or more *PIGA* gene mutations in their hematopoietic stem cells, which leads to abnormal blood cells. As the abnormal hematopoietic stem cells multiply, populations of abnormal blood cells are formed, alongside normal blood cells produced by normal hematopoietic stem cells. The proportion of abnormal blood cells in the body affects the severity of the signs and symptoms of paroxysmal nocturnal hemoglobinuria.

Researchers have identified more than 100 somatic mutations in the *PIGA* gene. Some of these mutations alter the numbers or types of protein building blocks (amino acids) in phosphatidylinositol glycan class A, which impair its function. Other mutations result in the insertion of a premature stop signal in the instructions for making phosphatidylinositol glycan class A. As a result, an abnormally small protein, which is usually unstable, is produced.

Chromosomal Location

Cytogenetic Location: Xp22.2, which is the short (p) arm of the X chromosome at position 22.2

Molecular Location: base pairs 15,319,451 to 15,335,554 on the X chromosome (Homo sapiens Annotation Release 108, GRCh38.p7) (NCBI)



Credit: Genome Decoration Page/NCBI

Other Names for This Gene

- GLCNAC-PI synthesis protein
- GPI anchor biosynthesis
- GPI3
- phosphatidylinositol glycan anchor biosynthesis, class A
- phosphatidylinositol glycan anchor biosynthesis, class A (paroxysmal nocturnal hemoglobinuria)
- phosphatidylinositol-glycan biosynthesis, class A protein
- phosphatidylinositol glycan, class A (paroxysmal nocturnal hemoglobinuria)
- phosphatidylinositol N-acetylglucosaminyltransferase subunit A
- PIG-A
- PIGA_HUMAN

Additional Information & Resources

Educational Resources

 Essentials of Glycobiology (first edition, 1999): Mutations in the GPI Anchor Biosynthetic Pathway https://www.ncbi.nlm.nih.gov/books/NBK20711/

Scientific Articles on PubMed

PubMed

https://www.ncbi.nlm.nih.gov/pubmed?term=%28PIGA%5BTIAB%5D%29+OR+%28%28GPI3%5BTIAB%5D%29+OR+%28PIG-A%5BTIAB%5D%29+OR+%28GPI+anchor+biosynthesis%5BTIAB%5D%29%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1440+days%22%5Bdp%5D

OMIM

 PHOSPHATIDYLINOSITOL GLYCAN ANCHOR BIOSYNTHESIS CLASS A PROTEIN

http://omim.org/entry/311770

Research Resources

- Atlas of Genetics and Cytogenetics in Oncology and Haematology http://atlasgeneticsoncology.org/Genes/GC_PIGA.html
- ClinVar https://www.ncbi.nlm.nih.gov/clinvar?term=PIGA%5Bgene%5D
- HGNC Gene Family: Glycosyl transferases group 1 domain containing http://www.genenames.org/cgi-bin/genefamilies/set/427
- HGNC Gene Family: Phosphatidylinositol glycan anchor biosynthesis http://www.genenames.org/cgi-bin/genefamilies/set/680
- HGNC Gene Symbol Report http://www.genenames.org/cgi-bin/gene_symbol_report?q=data/ hgnc_data.php&hgnc_id=8957
- NCBI Gene https://www.ncbi.nlm.nih.gov/gene/5277
- UniProt http://www.uniprot.org/uniprot/P37287

Sources for This Summary

- Brodsky RA, Hu R. PIG-A mutations in paroxysmal nocturnal hemoglobinuria and in normal hematopoiesis. Leuk Lymphoma. 2006 Jul;47(7):1215-21. Review.
 Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16923549
- Inoue N, Murakami Y, Kinoshita T. Molecular genetics of paroxysmal nocturnal hemoglobinuria. Int J Hematol. 2003 Feb;77(2):107-12. Review.
 Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/12627844

- Mortazavi Y, Merk B, McIntosh J, Marsh JC, Schrezenmeier H, Rutherford TR; BIOMED II
 Pathophysiology and Treatment of Aplastic Anaemia Study Group. The spectrum of PIG-A gene
 mutations in aplastic anemia/paroxysmal nocturnal hemoglobinuria (AA/PNH): a high incidence of
 multiple mutations and evidence of a mutational hot spot. Blood. 2003 Apr 1;101(7):2833-41. Epub
 2002 Nov 7.
 - Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/12424196
- OMIM: PHOSPHATIDYLINOSITOL GLYCAN ANCHOR BIOSYNTHESIS CLASS A PROTEIN http://omim.org/entry/311770
- Rosti V. The molecular basis of paroxysmal nocturnal hemoglobinuria. Haematologica. 2000 Jan; 85(1):82-7. Review.
 Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/10629597

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